

GenCore version 4.5  
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OM protein - protein search, using sw model

Run on: April 25, 2000, 20:10:14 ; Search time 85.2 Seconds  
(without alignments)  
176.812 Million cell updates/sec

Title: US-09-125-005-6  
Perfect score: 3384  
Sequence: 1 MAQSTATSPDGGTTFEHLWS.....PDCKARKQPIKEEFAEIH 636

Scoring table: BIOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 188963 seqs, 23686106 residues

Total number of hits satisfying chosen parameters: 188963

Minimum DB seq length: 0

Maximum DB seq length: 1000000

Post-processing: Minimum Match 0%

Listing first 45 summaries

Database : A\_Geneseq\_36.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

## SUMMARIES

Result No.	Query Match	Score	Length	DB ID	Description
1	3384	100.0	636	1 W36184	Human p53 tumour s
2	3367.5	99.5	635	1 W30661	Human NBS-1 alpha
3	3304.5	97.7	637	1 W36182	Monkey p53 tumour s
4	3330	92.5	588	1 W36189	Human p53 tumour s
5	3058	90.4	587	1 W36187	Human p53 tumour s
6	2830.5	83.6	589	1 W36185	Mouse p53 tumour s
7	2624	77.5	499	1 W36190	Human p53 tumour s
8	2574	76.1	499	1 W36183	Monkey p53 tumour s
9	2330.5	68.9	506	1 W36188	Human p53 tumour s
10	726.5	21.5	401	1 W28487	Human p53 protein
11	725.5	21.4	401	1 W28488	Human p53 protein
12	724.5	21.4	355	1 W13950	Del356-393 modifie
13	724.5	21.4	363	1 W13954	Modified p53 varia
14	724.5	21.4	393	1 R22238	Sequence of 53 KD
15	724.5	21.4	393	1 R26738	p53. Synthetic onc
16	724.5	21.4	393	1 R79658	Human p53 protein.
17	724.5	21.4	393	1 R94623	p53 protein. Recom
18	724.5	21.4	393	1 R91933	Wild type p53 prot
19	724.5	21.4	393	1 W02617	Human p53 tumour s
20	724.5	21.4	393	1 W05344	Human p53. New hum
21	724.5	21.4	393	1 W05348	Human wild-type p5
22	724.5	21.4	393	1 W57242	Human p53 protein
23	724.5	21.4	393	1 W57243	Human p53 protein
24	724.5	21.4	393	1 W48658	Amino acid sequenc
25	724.5	21.4	393	1 W69217	Human wild-type p5
26	724.5	21.4	393	1 W69718	Human p53 used in
27	724.5	21.4	393	1 Y03191	Amino acid sequenc
28	724.5	21.4	438	1 R74272	Tumour suppressor
29	724.5	21.4	533	1 W19763	p53-GM-CSF immuno
30	723.5	21.4	363	1 W13971	Modified p53 varia
31	723.5	21.4	393	1 W13949	T284R modified hum
32	723.5	21.4	393	1 W13953	T284K modified hum
33	723.5	21.4	393	1 W57244	Human p53 protein
34	723.5	21.4	393	1 W57245	Human p53 protein

35 723.5 21.4 393 1 W84270  
36 723.5 21.4 438 1 R50088  
37 720.5 21.3 363 1 W13972  
38 720.5 21.3 393 1 W05347  
39 720.5 21.3 393 1 W13951  
40 720 21.3 354 1 R51874  
41 719.5 21.3 363 1 W13973  
42 719.5 21.3 363 1 W13974  
43 719.5 21.3 393 1 W05345  
44 719.5 21.3 393 1 W05346  
45 719.5 21.3 393 1 W13968

Human p53 protein.  
p53 tumour suppress  
Modified p53 varia  
Human p53 mutant R  
Human tumour-deriv  
Human p53 amino ac  
Modified p53 varia  
Human p53 mutant N  
Human p53 mutant R  
Modified p53 varia

## ALIGNMENTS

RESULT 1

ID W36184 standard; Protein; 636 AA.

AC W36184; 1998 (first entry)

DE Human p53 tumour suppressor-related protein SR-p70a.

KW SR-p70; human; transcription factor; p53; tumour suppressor gene;

KW homology; differential splicing; diagnosis; cancer; neuroblastoma;

KW gene therapy; apoptosis.

OS Homo sapiens.

PN W09728186-A1.

PD 07-AUG-1997.

PF 03-FEB-1997; F00214.

PR 02-FEB-1996; FR-001309.

PA (SNFI ) SANOFI SA.

PI Caput D, Ferrara P, Kaghad AM;

DR WPI; 97-402550/37.

DR N-PSDB; V01498.

PT New polypeptide(s) encoded by the SR-p70 tumour suppressor gene -

PT and related nucleic acid, useful for diagnosis and treatment of

PT tumours

PS Claim 7: Fig 6: 136pp; French.

CC This is the amino acid sequence of the human protein SR-p70a. SR-p70

CC are transcription factors which may control the activity of p53-regulated

CC genes, and are expressed by tumour suppressor genes related to the p53

CC gene family. The gene sequence was isolated from the human colon

CC adenocarcinoma cell line Ht-29, using primers V01506-7. The sequence

CC can be used in the diagnosis and monitoring of cancer, especially

CC neuroblastoma. The nucleic acid sequences and corresponding antisense

CC sequences, are also useful in gene therapy, e.g. to regulate apoptosis.

SQ Sequence 636 AA;

Query Match

Best Local Similarity 100.0%; Score 3384; DB 1; Length 636;

Matches 636; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MAQSTATSPDGGTTFEHLWSLEPDSYFDLPQSSRGNNVEVGGTSSMDVFLHCGMTTS 60

Db 1 MAQSTATSPDGGTTFEHLWSLEPDSYFDLPQSSRGNNVEVGGTSSMDVFLHCGMTTS 60

QY 61 VMAQFNLLSTMDQMSRAASAPYTPHEAASVTPHSYAQPSSTFTDTSAPVIPSNTD 120

Db 61 VMAQFNLLSTMDQMSRAASAPYTPHEAASVTPHSYAQPSSTFTDTSAPVIPSNTD 120

QY 121 YPGPHFEVTFQSSSTAKSATWTYSPLLLKLYCQIAKTCPIQIKVSTPPPGTAIRAMPV 180

Db 121 YPGPHFEVTFQSSSTAKSATWTYSPLLLKLYCQIAKTCPIQIKVSTPPPGTAIRAMPV 180

QY 181 YKKAHVTDVVKPCNHELGRDENEGOSAPASHLIRVEGNLSQYVDDPVTGRQSVVVPY 240

Db 181 YKKAHVTDVVKPCNHELGRDENEGOSAPASHLIRVEGNLSQYVDDPVTGRQSVVVPY 240

QY 241 EPPQVGTEFTIILNFCNCSVCVGGMNRRLPILIIITLMDRGQVLGRSFEGRICACRGR 300

Db 241 EPPQVGTEFTIILNFCNCSVCVGGMNRRLPILIIITLMDRGQVLGRSFEGRICACRGR 300



Query Match 97.7%; Score 3304.5; DB 1; Length 637;  
Best Local Similarity 97.5%; Pred. No. 18-280;  
Matches 621; Conservative 4; Mismatches 11; Indels 1; Gaps 1;

QY 1 MAQSTATSPDGGTTFFHLWSSLEPDSYFDLPQSSRGNEVWVGDTSSMDVFLHGMITS 60  
DB 1 MAQSTTSPDGGTTFFHLWSSLEPDSYFDLPQSSRGNEVWVGDTSSMDVFLHGMITS 60

QY 61 VMAQFNLLSTMDQSSRAASAPYTPHEAASVPTHSPTAQSSTFTDMSAPVIPSNTD 120  
DB 61 VMAQFNLLSTMDQSSRAASAPYTPHEAASVPTHSPTAQSSTFTDMSAPVIPSNTD 120

QY 121 YPGPHFEVTFQSSSTAKSATWYSPLLKLYCQIAKTCPIQIKVSTPPPGTATRAMPV 180  
DB 121 YPGPHFEVTFQSSSTAKSATWYSPLLKLYCQIAKTCPIQIKVSTPPPGTATRAMPV 180

QY 181 YKKAHVTDVYKRCNHELGRDFNEGOSAPASHLIRVEGNNSQYVDDPVTGRQSVVY 240  
DB 181 YKKAHVTDVYKRCNHELGRDFNEGOSAPASHLIRVEGNNSQYVDDPVTGRQSVVY 240

QY 241 EPPQVGTETTYLYNFMCHSSCVGGMNRRPILIIITLEMRDGOVLGRSFEGRICACPR 300  
DB 241 EPPQVGTETTYLYNFMCHSSCVGGMNRRPILIIITLEMRDGOVLGRSFEGRICACPR 300

QY 301 DRKADEHYREOQALNESSAKNGAASKRAFQKSPAPVAPALGAGVYKRRRGGEDTYVLOVR 360  
DB 301 DRKADEHYREOQALNESSAKNGAASKRAFQKSPAPVAPALGAGVYKRRRGGEDTYVLOVR 360

QY 361 GRENFIILMKLESLELMELVPOPLVDSYRQOQLLQRPSSHLPQPSYGPVLSPMKNVHGG 420  
DB 361 GRENFIILMKLESLELMELVPOPLVDSYRQOQLLQRPSSHLPQPSYGPVLSPMKNVHGG 420

QY 421 MNKLPVNLVGGPPHSSAATPNLGPVGMNHNHGHAVPANGEMSSSHSAQSVSGSH 480  
DB 421 MNKLPVNLVGGPPHSSAATPNLGPVGMNHNHGHAVPANGEMSSSHSAQSVSGSH 480

QY 481 CTPPPYHADPSLVSLTGLGCPNCIEYFTSQGLQSIYHLQNTIEDLGALKIPEQYRMT 540  
DB 481 CTPPPYHADPSLVSLTGLGCPNCIEYFTSQGLQSIYHLQNTIEDLGALKIPEQYRMT 540

QY 541 IWRGLQDLKQGHY-STAQQLRSSNAATISIGSGELQQRVMEAVHFRVHTTIPNR 599  
DB 541 IWRGLQDLKQGHY-STAQQLRSSNAATISIGSGELQQRVMEAVHFRVHTTIPNR 600

QY 600 GPGGPGDEWADFGLPDCARKQPIKEEFTAEI 636  
DB 601 GPGGPGDEWADFGLPDCARKQPIKEEFTAEI 637

RESULT 4  
W36189  
ID W36189 standard; Protein: 588 AA.  
AC W36189;  
DT 27-APR-1998 (first entry)  
DE Human p53 tumour suppressor-related protein SR-p70f.  
KW SR-p70; human; transcription factor; p53; tumour suppressor gene;  
KW homology; differential splicing; diagnosis; cancer; neuroblastoma;  
KW gene therapy; apoptosis.  
OS Homo sapiens.  
PN WO9728186-A1.  
PD 07-AUG-1997.  
PF 03-FEB-1997; F00214.  
PR 02-FEB-1996; FR-001309.  
PA (SNEI) SANOI SA.  
PI Caput D, Ferrara P, Kaghad AM;  
DR WPI; 97-402550/37.  
DR N-PSDB; V01504.  
PT New polypeptide(s) encoded by the SR-p70 tumour suppressor gene -  
PT and related nucleic acid, useful for diagnosis and treatment of  
PT tumours  
PS Claim 1; Page 69-70; 136pp; French.  
CC This is the amino acid sequence of the human protein SR-p70f. SR-p70 are  
CC transcription factors which may control the activity of p53-regulated

CC genes, and are expressed by tumour suppressor genes related to the p53  
CC gene family. The gene sequence was isolated from the human neuroblastoma  
CC cell line SR-N-SH, using primers V01515 and V01518. The SR-p70f gene  
CC sequence contains a 98 bp deletion between bases 24-25 as compared to  
CC the SR-p70a sequence (V01498). This deletion causes a loss of the  
CC translation initiation codon found in SR-p70a, resulting in the use of  
CC a downstream AUG (corresponding to an internal Met codon in SR-p70a).  
CC The protein is truncated by 48 amino acids at the N-terminus as compared  
CC to the SR-p70a protein (W36184). The sequence can be used in the  
CC diagnosis and monitoring of cancer, especially neuroblastoma. The  
CC nucleic acid sequences and corresponding antisense sequences, are also  
CC useful in gene therapy, e.g. to regulate apoptosis.  
SQ Sequence 588 AA;

Query Match 92.5%; Score 3130; DB 1; Length 588;  
Best Local Similarity 100.0%; Pred. No. 1.7e-265;  
Matches 588; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 49 MDVPHLEGMTSVMAQFNLLSTMDQSSRAASAPYTPHEAASVPTHSPTAQSSTFTD 108  
DB 1 MDVPHLEGMTSVMAQFNLLSTMDQSSRAASAPYTPHEAASVPTHSPTAQSSTFTD 60

QY 109 MSPAPVIPSNTDYPGPHFEVTFQSSSTAKSATWYSPLLKLYCQIAKTCPIQIKVSTP 168  
DB 61 MSPAPVIPSNTDYPGPHFEVTFQSSSTAKSATWYSPLLKLYCQIAKTCPIQIKVSTP 120

QY 169 PPGTATRAMPVYKKAHVTDVYKRCNHELGRDFNEGOSAPASHLIRVEGNNSQYVDD 228  
DB 121 PPGTATRAMPVYKKAHVTDVYKRCNHELGRDFNEGOSAPASHLIRVEGNNSQYVDD 180

QY 229 PVTGRQSVVYVPPQVGTETTYLYNFMCHSSCVGGMNRRPILIIITLEMRDGOVLGR 288  
DB 181 PVTGRQSVVYVPPQVGTETTYLYNFMCHSSCVGGMNRRPILIIITLEMRDGOVLGR 240

QY 289 SFEGRICACPRDKADEHYREOQALNESSAKNGAASKRAFQKSPAPVAPALGAGVYKRR 348  
DB 241 SFEGRICACPRDKADEHYREOQALNESSAKNGAASKRAFQKSPAPVAPALGAGVYKRR 300

QY 349 HGDEDTYLYVGRGRENFEILMKLESLELMELVPOPLVDSYRQOQLLQRPSSHLPQPSY 408  
DB 301 HGDEDTYLYVGRGRENFEILMKLESLELMELVPOPLVDSYRQOQLLQRPSSHLPQPSY 360

QY 409 PVLSPMKNVHGGMKNLPSVNLVGGPPHSSAATPNLGPVGMNHNHGHAVPANGEMSS 468  
DB 361 PVLSPMKNVHGGMKNLPSVNLVGGPPHSSAATPNLGPVGMNHNHGHAVPANGEMSS 420

QY 469 SHSAQSVSGSHCTPPPYHADPSLVSLTGLGCPNCIEYFTSQGLQSIYHLQNTIEDL 528  
DB 421 SHSAQSVSGSHCTPPPYHADPSLVSLTGLGCPNCIEYFTSQGLQSIYHLQNTIEDL 480

QY 529 GALKIPEQYRMTIWRGLQDLKQGHY-STAQQLRSSNAATISIGSGELQQRVMEAVH 588  
DB 481 GALKIPEQYRMTIWRGLQDLKQGHY-STAQQLRSSNAATISIGSGELQQRVMEAVH 540

QY 589 RVHRTTIPNRGGPGGDEWADFGLPDCARKQPIKEEFTAEI 636  
DB 541 RVHRTTIPNRGGPGGDEWADFGLPDCARKQPIKEEFTAEI 588

RESULT 5  
W36187  
ID W36187 standard; Protein: 587 AA.  
AC W36187;  
DT 27-APR-1998 (first entry)  
DE Human p53 tumour suppressor-related protein SR-p70d.  
KW SR-p70; human; transcription factor; p53; tumour suppressor gene;  
KW homology; differential splicing; diagnosis; cancer; neuroblastoma;  
KW gene therapy; apoptosis.  
OS Homo sapiens.  
PN WO9728186-A1.  
PD 07-AUG-1997.  
PF 03-FEB-1997; F00214.

02-FEB-1996; FR-001309.  
 (SNFI) SANOFI SA.  
 Caput D, Ferrara P, Kaghad AM;  
 WPI: 97-402550/37.  
 N-PSDB; V01502.  
 New polypeptide(s) encoded by the SR-p70 tumour suppressor gene -  
 and related nucleic acid, useful for diagnosis and treatment of  
 tumours  
 Claim 1; Page 62-64; 136pp; French.  
 This is the amino acid sequence of the human protein SR-p70d. R-p70 are  
 transcription factors which may control the activity of p53-regulated  
 genes, and are expressed by tumour suppressor genes related to the p53  
 gene family. The gene sequence was isolated from the human colon  
 neuroblastoma cell line IMR-32, using primers V01512-13. The SR-p70d  
 protein sequence is 49 amino acids shorter with a divergence of the  
 first 13 amino acids as compared to the SR-p70a protein (W36184).  
 The sequence can be used in the diagnosis and monitoring of cancer,  
 especially neuroblastoma. The nucleic acid sequences and corresponding  
 antisense sequences, are also useful in gene therapy, e.g. to regulate  
 apoptosis.  
 Sequence 587 AA;

Query Match 90.4%; Score 3058; DB 1; Length 587;  
 Best Local Similarity 100.0%; Pred. No. 3.3e-259;  
 Matches 574; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

63 AOFNLLSSTMDQMSRAASAPYTPHEAASVTPHSPVAQPSSTEDTMSAPVPSNTDYP 122  
 14 AOFNLLSSTMDQMSRAASAPYTPHEAASVTPHSPVAQPSSTEDTMSAPVPSNTDYP 73  
 123 GPHFEVTFQOSTAKSATWTYSPLKLYCOIAKTCPIQIKVSTPPPGTAIRAMPVYK 182  
 74 GPHFEVTFQOSTAKSATWTYSPLKLYCOIAKTCPIQIKVSTPPPGTAIRAMPVYK 133  
 183 KAEHTVDVVKCPNHELGRDNEGOSAPASHLIRVEGNLSQYVDDPVTGRQSVVYP 242  
 134 KAEHTVDVVKCPNHELGRDNEGOSAPASHLIRVEGNLSQYVDDPVTGRQSVVYP 193  
 243 POGVTEFTILYFNCNSCVGMNRRPILIIITLMDRGQVLRGRSFEGRICACGRDR 302  
 194 POGVTEFTILYFNCNSCVGMNRRPILIIITLMDRGQVLRGRSFEGRICACGRDR 253  
 303 KADEHYREQQALNESSAKRAFKQSPAPALGAGVKKRRHGDEDTYVLYQVGR 362  
 254 KADEHYREQQALNESSAKRAFKQSPAPALGAGVKKRRHGDEDTYVLYQVGR 313  
 363 ENFEILMKLSLELMELVPOPLVDSYRQOQQLLQRPSPHLPSPYGPVLSPMKRVHGMN 422  
 314 ENFEILMKLSLELMELVPOPLVDSYRQOQQLLQRPSPHLPSPYGPVLSPMKRVHGMN 373  
 423 KLPSYNQLVGPPPHSSAATPNLGPVGMNHNHGHAVPANGEMSSSHSAQSMVSGSHCT 482  
 374 KLPSYNQLVGPPPHSSAATPNLGPVGMNHNHGHAVPANGEMSSSHSAQSMVSGSHCT 433  
 483 PPPPHADPSLVFTGLGCPNCIEYFTSQGLSIYHLQNTIEDLGALEKIPQYRMTIW 542  
 434 PPPPHADPSLVFTGLGCPNCIEYFTSQGLSIYHLQNTIEDLGALEKIPQYRMTIW 493  
 543 RLQDLKQGHDIYTAQQLLRSSNAATISIGGSGELQQRVMEAVHFRVHTITIPNRRGP 602  
 494 RLQDLKQGHDIYTAQQLLRSSNAATISIGGSGELQQRVMEAVHFRVHTITIPNRRGP 553  
 603 GGGPDWADFGDLPDCKARKQPIKEEFTAEIH 636  
 554 GGGPDWADFGDLPDCKARKQPIKEEFTAEIH 587

RESULT 6  
 36185  
 C W36185 standard; Protein; 589 AA.  
 C W36185;  
 T 27-APR-1998 (first entry)

DE Mouse p53 tumour suppressor-related protein SR-p70c.  
 KW SR-p70; mouse; transcription factor; p53; tumour suppressor gene;  
 KW homology; differential splicing; diagnosis; cancer; neuroblastoma;  
 KW gene therapy; apoptosis.  
 OS Mus musculus.  
 PN W09728186-A1.  
 PD 07-AUG-1997.  
 PF 03-FEB-1997; F00214.  
 PR 03-FEB-1996; FR-001309.  
 PA (SNFI) SANOFI SA.  
 PI Caput D, Ferrara P, Kaghad AM;  
 DR WPI: 97-402550/37.  
 N-PSDB; V01499.  
 PT New polypeptide(s) encoded by the SR-p70 tumour suppressor gene -  
 and related nucleic acid, useful for diagnosis and treatment of  
 tumours  
 Claim 1; Fig 7; 136pp; French.  
 This is the amino acid sequence of the mouse protein SR-p70c. SR-p70  
 are transcription factors which may control the activity of p53-regulated  
 genes, and are expressed by tumour suppressor genes related to the p53  
 gene family. The gene sequence was isolated from the mouse pituitary  
 tumour cell line Atf-20, using primers V01508-9. The sequence can be  
 used in the diagnosis and monitoring of cancer, especially neuroblastoma.  
 The nucleic acid sequences and corresponding antisense sequences, are  
 also useful in gene therapy, e.g. to regulate apoptosis.  
 Sequence 589 AA;

Query Match 83.6%; Score 2830.5; DB 1; Length 589;  
 Best Local Similarity 91.7%; Pred. No. 2.7e-239;  
 Matches 531; Conservative 19; Mismatches 22; Indels 7; Gaps 4;

63 AOFNLLSSTMDQMSRAASAPYTPHEAASVTPHSPVAQPSSTEDTMSAPVPSNTDYP 122  
 13 AOFNLLSSTMDQMSRAASAPYTPHEAASVTPHSPVAQPSSTEDTMSAPVPSNTDYP 72  
 123 GPHFEVTFQOSTAKSATWTYSPLKLYCOIAKTCPIQIKVSTPPPGTAIRAMPVYK 182  
 73 GPHFEVTFQOSTAKSATWTYSPLKLYCOIAKTCPIQIKVSTPPPGTAIRAMPVYK 132  
 183 KAEHTVDVVKCPNHELGRDNEGOSAPASHLIRVEGNLSQYVDDPVTGRQSVVYP 242  
 133 KAEHTVDVVKCPNHELGRDNEGOSAPASHLIRVEGNLSQYVDDPVTGRQSVVYP 192  
 243 POGVTEFTILYFNCNSCVGMNRRPILIIITLMDRGQVLRGRSFEGRICACGRDR 302  
 193 POGVTEFTILYFNCNSCVGMNRRPILIIITLMDRGQVLRGRSFEGRICACGRDR 252  
 303 KADEHYREQQALNESSAKRAFKQSPAPALGAGVKKRRHGDEDTYVLYQVGR 362  
 253 KADEHYREQQALNESSAKRAFKQSPAPALGAGVKKRRHGDEDTYVLYQVGR 312  
 363 ENFEILMKLSLELMELVPOPLVDSYRQOQQLLQRPSPHLPSPYGPVLSPMKRVHGMN 420  
 313 ENFEILMKLSLELMELVPOPLVDSYRQOQQLLQRPSPHLPSPYGPVLSPMKRVHGMN 372  
 421 MNKLPVSNQLVGPPPHSSAATPNLGPVGMNHNHGHAVPANGEMSSSHSAQSMVSGSH 480  
 373 VNKLPVSNQLVGPPPHSSAAGPNLGPNGSGLNHSMPANGEMNGHSSOTMVSGSH 432  
 481 CTPPPPHADPSLVFTGLGCPNCIEYFTSQGLSIYHLQNTIEDLGALEKIPQYRMT 540  
 433 CTPPPPHADPSLVFTGLGCPNCIEFTSQGLSIYHLQNTIEDLGALEKIPQYRMT 492  
 541 IWRGLQDLKQGHDIYTAQQLLRSSNAATISIGGSGELQQRVMEAVHFRVHTITIPNR 599  
 493 IWRGLQDLKQGHDIYTAQQLLRSSNAATISIGGSGELQQRVMEAVHFRVHTITIPNR 550  
 600 GGGPDWADFGDLPDCKARKQPIKEEFTAEIH 636  
 551 GGGPDWADFGDLPDCKARKQPIKEEFTAEIH 589

RESULT 7  
W36190  
ID W36190 standard; Protein; 499 AA.  
AC W36190;  
DC 27-APR-1998 (first entry)  
DE Human p53 tumour suppressor-related protein SR-p70b.  
KW SR-p70; human; transcription factor; p53; tumour suppressor gene;  
KW homology; differential splicing; diagnosis; cancer; neuroblastoma;  
KW gene therapy; apoptosis.  
OS Homo sapiens.  
PN WO9728186-A1.  
PD 07-AUG-1997.  
PF 03-FEB-1997; F00214.  
PR 02-FEB-1996; FR-001309.  
PA (SNFI) SANOFI SA.  
PI Caput D, Ferrara P, Kaghad AM;  
DR WPI; 97-402550/37.  
DR N-PSDB; V01505.  
PT New polypeptide(s) encoded by the SR-p70 tumour suppressor gene -  
and related nucleic acid, useful for diagnosis and treatment of  
tumours.  
PS Claim 1; page 72-73; 136pp; French.  
CC This is the amino acid sequence of the human protein SR-p70b. SR-p70 are  
transcription factors which may control the activity of p53-regulated  
genes, and are expressed by tumour suppressor genes related to the p53  
gene family. The gene sequence was isolated from the human neuroblastoma  
cell line SK-N-SH, using primers V01515 and V01518. The SR-p70b gene  
sequence contains a 94 bp deletion between bases 1516-1517 as compared  
to the SR-p70a sequence (V01498). This deletion causes a reading frame  
shift resulting in the generation of a stop codon at position 1498-1500.  
The resultant protein is truncated by 137 amino acids as compared to the  
SR-p70a protein (W36184). The sequence can be used in the diagnosis and  
monitoring of cancer, especially neuroblastoma. The nucleic acid  
sequences and corresponding antisense sequences, are also useful in gene  
therapy, e.g. to regulate apoptosis.  
SQ Sequence 499 AA;

Query Match 77.5%; Score 2624; DB 1; Length 499;  
Best Local Similarity 100.0%; Pred. No. 2.5e-221;  
Matches 494; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 1 MAQSTATSPDGGTTFEHLWSSLEPDSYFDLPQSSRGNNVVGTTDSSMDVFLHGMGTTTS 60  
DB 1 MAQSTATSPDGGTTFEHLWSSLEPDSYFDLPQSSRGNNVVGTTDSSMDVFLHGMGTTTS 60  
QY 61 VMAQFNLLSSTMDQSSRAASASPYTPEHAASVTHSPYAQPSSTFTDMSAPVIPSNTD 120  
DB 61 VMAQFNLLSSTMDQSSRAASASPYTPEHAASVTHSPYAQPSSTFTDMSAPVIPSNTD 120  
QY 121 YPGPHFEVTFQOSTAKSATWTYSPLLKLYCOIAKTCPIQIKVSTPPPGTAIRAMPV 180  
DB 121 YPGPHFEVTFQOSTAKSATWTYSPLLKLYCOIAKTCPIQIKVSTPPPGTAIRAMPV 180  
QY 181 YKKAHEVTDVVKRCNPHELGRDFNEGOSAPASHLIRVEGNLSQYVDDPVTGROSVVVPY 240  
DB 181 YKKAHEVTDVVKRCNPHELGRDFNEGOSAPASHLIRVEGNLSQYVDDPVTGROSVVVPY 240  
QY 241 EPPQVGTFTILYFNMCNCSVCGMNRRLIITILEMRDQGVLGRRSFEGRICACPGR 300  
DB 241 EPPQVGTFTILYFNMCNCSVCGMNRRLIITILEMRDQGVLGRRSFEGRICACPGR 300  
QY 301 DRKADEHYRQOALNESSAKNGAASKRAFKQSPAPALGAGVKRRHGDDETYIYQVR 360  
DB 301 DRKADEHYRQOALNESSAKNGAASKRAFKQSPAPALGAGVKRRHGDDETYIYQVR 360  
QY 361 GRENFELMKLESLELMELVPQPLVDSYRQOQLLRPSHLQPPSYGVLSPMNKHYGG 420  
DB 361 GRENFELMKLESLELMELVPQPLVDSYRQOQLLRPSHLQPPSYGVLSPMNKHYGG 420  
QY 421 MNKLPVSNQVLGQPPPHSSAATPNLGPVPGMKNHGHAVPANGEMSSSSHAQSMVSGSH 480  
DB 421 MNKLPVSNQVLGQPPPHSSAATPNLGPVPGMKNHGHAVPANGEMSSSSHAQSMVSGSH 480

QY 481 CTPPPPPYHADPSLV 494  
DB 481 CTPPPPPYHADPSLV 494  
RESULT 8  
W36183  
ID W36183 standard; Protein; 499 AA.  
AC W36183;  
DC 27-APR-1998 (first entry)  
DE Monkey p53 tumour suppressor-related protein SR-p70b.  
KW SR-p70; monkey; transcription factor; p53; tumour suppressor gene;  
KW homology; differential splicing; diagnosis; cancer; neuroblastoma;  
KW gene therapy; apoptosis.  
OS Cercopithecus aethiops.  
PN WO9728186-A1.  
PD 07-AUG-1997.  
PF 03-FEB-1997; F00214.  
PR 02-FEB-1996; FR-001309.  
PA (SNFI) SANOFI SA.  
PI Caput D, Ferrara P, Kaghad AM;  
DR WPI; 97-402550/37.  
DR N-PSDB; V01497.  
PT New polypeptide(s) encoded by the SR-p70 tumour suppressor gene -  
and related nucleic acid, useful for diagnosis and treatment of  
tumours.  
PS Claim 1; Fig 5; 136pp; French.  
CC This is the amino acid sequence of the protein SR-p70b from monkey cells.  
SR-p70 are transcription factors which may control the activity of  
p53-regulated genes, and are expressed by tumour suppressor genes related  
to the p53 gene family. The gene sequence was isolated from a cDNA  
library by sequencing the inserts and comparing to sequence databases.  
The protein sequence contains regions of homology to the p53 protein.  
The SR-p70b gene sequence was isolated simultaneously with the SR-p70a  
sequence (V01496) from the library and is created by differential  
splicing of the SR-p70 mRNA sequence. The sequences can be used in the  
diagnosis and monitoring of cancer, especially neuroblastoma. The  
nucleic acid sequences and corresponding antisense sequences, are also  
useful in gene therapy, e.g. to regulate apoptosis.  
SQ Sequence 499 AA;

Query Match 76.1%; Score 2574; DB 1; Length 499;  
Best Local Similarity 97.8%; Pred. No. 5.8e-217;  
Matches 483; Conservative 3; Mismatches 8; Indels 0; Gaps 0;  
QY 1 MAQSTATSPDGGTTFEHLWSSLEPDSYFDLPQSSRGNNVVGTTDSSMDVFLHGMGTTTS 60  
DB 1 MAQSTATSPDGGTTFEHLWSSLEPDSYFDLPQSSRGNNVVGTTDSSMDVFLHGMGTTTS 60  
QY 61 VMAQFNLLSSTMDQSSRAASASPYTPEHAASVTHSPYAQPSSTFTDMSAPVIPSNTD 120  
DB 61 VMAQFNLLSSTMDQSSRAASASPYTPEHAASVTHSPYAQPSSTFTDMSAPVIPSNTD 120  
QY 121 YPGPHFEVTFQOSTAKSATWTYSPLLKLYCOIAKTCPIQIKVSTPPPGTAIRAMPV 180  
DB 121 YPGPHFEVTFQOSTAKSATWTYSPLLKLYCOIAKTCPIQIKVSTPPPGTAIRAMPV 180  
QY 181 YKKAHEVTDVVKRCNPHELGRDFNEGOSAPASHLIRVEGNLSQYVDDPVTGROSVVVPY 240  
DB 181 YKKAHEVTDVVKRCNPHELGRDFNEGOSAPASHLIRVEGNLSQYVDDPVTGROSVVVPY 240  
QY 241 EPPQVGTFTILYFNMCNCSVCGMNRRLIITILEMRDQGVLGRRSFEGRICACPGR 300  
DB 241 EPPQVGTFTILYFNMCNCSVCGMNRRLIITILEMRDQGVLGRRSFEGRICACPGR 300  
QY 301 DRKADEHYRQOALNESSAKNGAASKRAFKQSPAPALGAGVKRRHGDDETYIYQVR 360  
DB 301 DRKADEHYRQOALNESSAKNGAASKRAFKQSPAPALGAGVKRRHGDDETYIYQVR 360  
QY 361 GRENFELMKLESLELMELVPQPLVDSYRQOQLLRPSHLQPPSYGVLSPMNKHYGG 420  
DB 361 GRENFELMKLESLELMELVPQPLVDSYRQOQLLRPSHLQPPSYGVLSPMNKHYGG 420

361 GRENFEILMKLSLELMELVPOPLVDYSYRQOQLLQRPSPHLPQPSYGPVLSPMKNVHGG 420  
421 MNKLPVNLVQGGPPHSSAATNLGPGVGMNLNRRHGHAVPANGEMSSSHSAQSMVSGSH 480  
422 VNKLPVNLVQGGPPHSSAATNLGPGVGMNLNRRHGHAVPANGEMSSSHSAQSMVSGSH 480  
481 CTPPPPYHADPSLV 494  
482 CTPPPPYHADPSLV 494  
RESULT 9  
36188  
W36188 standard; Protein; 506 AA.  
W36188; (first entry)  
Human p53 tumour suppressor-related protein SR-p70e.  
SR-p70; human; transcription factor; p53; tumour suppressor gene;  
homology; differential splicing; diagnosis; cancer; neuroblastoma;  
gene therapy; apoptosis.  
Homo sapiens.  
W09728186-AL.  
07-AUG-1997.  
03-FEB-1997; F00214.  
02-FEB-1996; FR-001309.  
(SNFI) SANOFI SA.  
Caput D, Ferrara P, Kaghad AM;  
WFI; 97-403550/37.  
N-PSDB; V01503.  
New polypeptide(s) encoded by the SR-p70 tumour suppressor gene -  
and related nucleic acid, useful for diagnosis and treatment of  
tumours  
Claim 1; Page 65-66; 136pp; French.  
This is the amino acid sequence of the human protein SR-p70e. SR-p70 are  
transcription factors which may control the activity of p53-regulated  
genes, and are expressed by tumour suppressor genes related to the p53  
gene family. The gene sequence was isolated from the human neuroblastoma  
cell line IMR-32, using primers V01515 and V01518. The SR-p70e gene  
sequence is identical to the SR-p70d sequence (V01502) except for 2  
deletions of 149 bp and 94 bp between bases 1049-1050 and 1188-1189  
respectively. The encoded protein is 49 amino acids shorter with a  
divergence of the first 13 amino acids and a divergence between amino  
acids 350-397 as compared to the SR-p70a protein (W36184). The sequence  
can be used in the diagnosis and monitoring of cancer, especially  
neuroblastoma. The nucleic acid sequences and corresponding antisense  
sequences, are also useful in gene therapy, e.g. to regulate apoptosis.  
Sequence 506 AA;  
Query Match 68.9%; Score 2330.5; DB 1; Length 506;  
Best Local Similarity 80.7%; Pred. No. 1.2e-195;  
Matches 464; Conservative 1; Mismatches 27; Indels 83; Gaps 6;  
63 AQNLSSNDQMSRAASASPTPEHAASVTHSPYAPQSPSTFTDMSAPVPSNTDYP 122  
14 AQNLSSNDQMSRAASASPTPEHAASVTHSPYAPQSPSTFTDMSAPVPSNTDYP 73  
123 GPHEFVTFQOSTAKSATWYSPLKLYLCQIAKTCPIQIKVSTPPPPGTATRAAMPYK 182  
74 GPHEFVTFQOSTAKSATWYSPLKLYLCQIAKTCPIQIKVSTPPPPGTATRAAMPYK 133  
183 KAEHVTDVVKRCNHELGRDFNEGOSAPASHLIRVEGNLSQYDDPVTGRQSVVVPYEP 242  
134 KAEHVTDVVKRCNHELGRDFNEGOSAPASHLIRVEGNLSQYDDPVTGRQSVVVPYEP 193  
243 POGTETFTILYNMNCSSCGMNRRLIITILEMRDQVILGRSFEGRICACPGDR 302  
194 POGTETFTILYNMNCSSCGMNRRLIITILEMRDQVILGRSFEGRICACPGDR 253  
303 KADEHYREQQALNESSAKNGAASRAFAKQSPAPVAPALGAGYKRRHGDDETYILOVRGR 362  
254 KADEHYREQQALNESSAKNGAASRAFAKQSPAPVAPALGAGYKRRHGDDETYILOVRGR 313

363 ENFEILMKLSLELMELVPOPLVDYSYRQOQLLQRPSPHLPQPSYGPVLSPMKNVHGGMN 422  
314 ENFEILMKLSLELMELVPOPLVDYSYRQOQLLQRP----- 351  
423 KLPVNLVQGGPPHSSAATNLGPGVGMNLNRRHGHAVPANGEMSSSHSAQSMVSGSHC 481  
351 --PRDAQ--QPWPRASQRDEQOQORP-----VHGLGVPL-----HS 384  
482 TTPPPPYHADPSLVSLFTGLGCPNCIEYFTSQGLQSIYHLQNLTIEDLGALKIPEQYRMTI 541  
385 ATPLPRRPQPR-----QDLGALKIPEQYRMTI 411  
542 WRGLDLKQGHDSYTAQQLRSSNAATISIGSGSELQORQVMEAVHFRVHTTITINRGG 601  
412 WRGLDLKQGHDSYTAQQLRSSNAATISIGSGSELQORQVMEAVHFRVHTTITINRGG 471  
602 PGGPDEWADFGFDLPCKARKQPIKEEFTAEIHH 636  
472 PGGPDEWADFGFDLPCKARKQPIKEEFTAEIHH 506  
RESULT 10  
W28487  
ID W28487 standard; Protein; 401 AA.  
AC W28487;  
DT 25-NOV-1997 (first entry)  
DE Human p53 protein variant V-393 encoded by p53145.  
KW Leucine zipper domain; LZP; oligomerisation domain; mutant; mutein;  
KW substitution; replacement; transactivation; viral protein VP16; HSV;  
KW anti-oncogene; hyperproliferation; cancer; restenosis;  
KW tumour suppression; apoptosis.  
OS Chimeric - Homo sapiens.  
OS Chimeric - Herpes simplex virus.  
OS Synthetic.  
PN W09704092-AL.  
PD 06-FEB-1997.  
PF 17-JUL-1996; F01111.  
PR 13-JUL-1995; FR-008729.  
PA (RHON) RHONE POULENC RORER SA.  
PI Bracco L, Conseiller E;  
DR WFI; 97-132633/12.  
DR N-PSDB; T86219.  
DR New p53 variants e.g. with oligomerisation domain replaced by  
PT leucine zipper - useful for treating hyper-proliferative disorders,  
PT esp. cancer and restenosis  
PS Claim 34; Pages 83-85; 133pp; French.  
CC Claimed variants of protein p53 have at least part of the p53  
transactivation domain (amino acids 1-74) Deleted and replaced by  
CC the transactivating domain (TD) from herpes simplex virus viral  
CC protein VP16 (amino acids 411-490). The present sequence is that of  
CC a specifically claimed p53 variant designated V-393 and comprising  
CC the VP16 TD with amino acids 75-393 of human wild-type p53. The p53  
CC variants are more active and more stable tumour suppressors and  
CC apoptosis-inducing agents than wild-type p53 and are active where  
CC the wild-type protein is not.  
SQ Sequence 401 AA;  
Query Match 21.5%; Score 726.5; DB 1; Length 401;  
Best Local Similarity 44.1%; Pred. No. 1.9e-55;  
Matches 162; Conservative 56; Mismatches 99; Indels 55; Gaps 11;  
29 FDLPSQSRGNEVVGTT--DSS-----MDVFHLEGMTTSVMAQFNLLSSTMDQSSRAA 80  
33 FDLMLGDGDSGPGGTFPHDSAPYGLADNADFEFQMTDALG-----IDYGGRA 84  
81 SASPYTPEHAASVTHSPYAPQSPSTFTDMSAPVPSNTDYPGPHEFVTFQOSTAKSA 140  
85 PAAP-TP-----AAPAPAPSWPLSSS-----VPSQKTYQSGYGRFLGFLHSQAKSV 130  
141 TWYTSPLKLYLCQIAKTCPIQIKVSTPPPPGTATRAAMPYKAEHVTDVVKRCNHEL 200  
131 TCTYSPALNKMFCQLAKTCVQLWYDSTPPPGTTRVRAAIYKQSQHMTVEVRRCPHERC 190





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1b 228 DCTTHNYMNCSCMGNNRRPILITLSDSGNLGRNSFEVRCACPGDRDRTEEE 287
2y 308 HYREQOALNESSAKNG-----AASKRAFKQSPAPVAGVKKRRHGHDEYVYLOV 359
1b 288 NLR-----KKGEPHHELPPGSTRALPNTSSSPQ-----PKKKPLDGEYFTLQI 332
2y 360 RGRENFEILMKLESLELME 379
1b 333 RGRERFEMFRELNEALELKD 352

RESULT 13
ID W13954 standard; Protein; 363 AA.
AC W13954;
DE Modified p53 variant (first entry)
KW p53; tumour suppressor; cancer; therapy; cell proliferation;
KW apoptosis; protein engineering; DNA binding.
CS Synthetic.
PN W09710843-A1.
PF 20-SEP-1996; U15188.
PR 22-SEP-1995; US-004802.
PR 21-AUG-1996; US-697221.
PR (WIST-) WISTAR INST ANATOMY & BIOLOGY.
PI Halazometis TD;
DR WPI; 97-203618/18.
PT R284K modified p53 protein having DNA binding ability - useful in
PT treatment of cancer
PT Example 1; 49-51; 82pp; English.
CC A modified p53 variant (W13954) comprises wild-type p53 (see
CC also W13948) having a deletion of the C-terminal 30 amino acids,
CC and is obtd. by site-directed mutagenesis of p53 DNA. Deletion of
CC the p53 C-terminal 30 amino acids activates the DNA binding of
CC common Class I p53 mutants (see also W13951-52). Novel modified
CC p53 variants (W13949-50, W13953-54, W13968-77), some contg.
CC C-terminal deletions, provide the means for pharmacological rescue
CC of p53 function in cancer patients. Nucleic acids coding for
CC modified p53 can be used for cancer gene therapy.
CC Sequence 363 AA;

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Query Match 21.4%; Score 724.5; DB 1; Length 363;
Best Local Similarity 42.1%; Pred. No. 2.5e-55;
Matches 160; Conservative 58; Mismatches 103; Indels 59; Gaps 10;

2y 14 TFEHLWSSLEPSTYFDLPQSSRGNNVGVGTDSSMDVFLHLEGMTTSVMAQFNLLSSMD 73
1b 18 TFSDLWKLLENVLSPLP-----SQAMDDLMLSPDD-----IEQWTFEDPGPD 61
2y 74 QMSRAASAPYTPHEAASVPTSHSPYAPQSSSTFDI-MSPAPV-----VIPSNTDYPGPHF 127
1b 62 EAPRMEFAAPPVAPAPATP-----AAP-----APAPSWPLSSVPSQKTTQSGVGF 109
2y 128 EVTFQOSSTAKSATWTYSPLLKLYCQIAKTCPIQIKVSTPPPGTAIRAMPVYKAEHV 187
1b 110 RLGLFHSHTAKSVTCYSPALNKMFCOLAKTCVQLWVDSTPPGTRVRAMAIYKQSHM 169
2y 188 TDVVKCPNHELGRDNFNEGQSPASHLIRVEGNNSQYVDDPVTGROSVVYVPEPQVGT 247
1b 170 TEVVRCPHHERGSD-SDG-LAPPOHLIRVEGNLRVEYLDLDRNTFRHSVVVYVPEPVS 227
2y 248 EFTTILYFMNCSSCVGGMNRRPILITILEMRDGOVLGRRRFEGRICACPGDRKADAED 307
1b 228 DCTTHNYMNCSCMGNNRRPILITLSDSGNLGRNSFEVRCACPGDRDRTEEE 287
2y 308 HYREQOALNESSAKNG-----AASKRAFKQSPAPVAGVKKRRHGHDEYVYLOV 359
1b 288 NLR-----KKGEPHHELPPGSTRALPNTSSSPQ-----PKKKPLDGEYFTLQI 332
2y 360 RGRENFEILMKLESLELME 379

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1b 333 RGRERFEMFRELNEALELKD 352
2y 333 RGRERFEMFRELNEALELKD 352

RESULT 14
ID R22238 standard; Protein; 393 AA.
AC R22238;
DE 23-JUL-1992 (first entry)
KW Sequence of 53 kD cellular protein.
KW Cancer therapy; cancer suppressor gene; oncogenesis.
OS Homo sapiens.
PN EP-475623-A.
PD 18-MAR-1992.
PF 23-AUG-1991; 307791.
PR 24-AUG-1990; US-573405.
PR (REGC) UNIV OF CALIFORNIA.
PI Lee WH, Chen PL;
DR WPI; 92-090221/12.
DR N-PSDB; Q22995.
PT Cloned p53 cDNA and protein prods. - for suppression of
PT neoplastic phenotype e.g. in osteo-sarcoma(s), leukaemia(s),
PT lymphoma(s), etc. English.
PS Claim 2; Page 14; 25pp; English.
CC p53 cDNA, or its gene prods., can be used to suppress and eradicate
CC cancers caused by defective, mutant or absent cancer suppressor
CC genes. Variant forms of p53 are found in human breast, lung or
CC colon carcinoma, lymphoma, leukaemia, etc., suggesting that mutation
CC of the p53 genes is involved in oncogenesis. Specifically 273 Arg
CC is replaced by 273 His, a mutation found exclusively in tumour cells.
CC Sequence 393 AA;

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Query Match 21.4%; Score 724.5; DB 1; Length 393;
Best Local Similarity 41.0%; Pred. No. 2.8e-55;
Matches 157; Conservative 57; Mismatches 104; Indels 65; Gaps 9;

QY 14 TFEHLWSSLEPSTYFDLPQSSRGNNVGVGTDSSMDVFLHLEGMTTSVMAQFNLLSSMD 73
1b 18 TFSDLWKLLENVLSPLP-----SQAMDDLMLSPDDIE 51
QY 74 QMSRAASAPYTPHEAASVPTSHSPYAPQSSSTFDI-MSPAPV-----IPSNTDYPGP 124
1b 52 QWTFEDPG-----PDEAPRMEFAAPPVAPAPATP-----PAPAPSWPLSSVPSQKTYGS 106
QY 125 HFEVTFQOSSTAKSATWTYSPLLKLYCQIAKTCPIQIKVSTPPPGTAIRAMPVYKKA 184
1b 107 YGRLGFLHSGTAKSVTCYSPALNKMFCOLAKTCVQLWVDSTPPGTRVRAMAIYKQS 166
QY 185 EHVTDVVKCPNHELGRDNFNEGQSPASHLIRVEGNNSQYVDDPVTGROSVVYVPEPQ 244
1b 167 QHMTVEVVRCPHHERGSD-SDG-LAPPOHLIRVEGNLRVEYLDLDRNTFRHSVVVYVPEPE 224
QY 245 VGTFTTILYFMNCSSCVGGMNRRPILITILEMRDGOVLGRRRFEGRICACPGDRKA 304
1b 225 VGSDCITTHNYMNCSCMGNNRRPILITILESSGNLGRNSFEVRCACPGDRRT 284
QY 305 DEDHYREQOALNESSAKNG-----AASKRAFKQSPAPVAGVKKRRHGHDEYTY 356
1b 285 EENLR-----KKGEPHHELPPGSTRALPNTSSSPQ-----PKKKPLDGEYFT 329
QY 357 LQVRGRENFEILMKLESLELME 379
1b 330 LQIRGRERFEMFRELNEALELKD 352

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RESULT 15
ID R26758 standard; peptide; 393 AA.
AC R26758;
DE 09-FEB-1993 (first entry)
DE p53.
KW Point mutation; translocation; proto-oncogene; cancer;

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KW antigen-presenting cell; T-cell; HLA.  
 OS Synthetic.  
 FH Key Location/Qualifiers  
 FT misc\_difference 273  
 FT /label= mutation  
 FT /note= "Arg -> any amino acid except Arg, see CC"  
 PN GB2253211-A.  
 PD 02-SEP-1992.  
 PR 26-FEB-1992; 004098.  
 PR 26-FEB-1991; GB-003974.  
 PA (NHSD ) NORSK HYDRO AS.  
 PI Eriksen JA, Gaudernack G, Geddedahl T;  
 DR WPI: 92-294575/36.  
 PT Synthetic oncogene-protein peptide - used for treating and  
 PT vaccinating against cancers  
 PS Disclosure; Page 10; 78pp; English.  
 CC New peptides, which have a point mutation or translocation compared to  
 CC the corresp. fragment of the proto-oncogene prod., correspond to,  
 CC completely cover or are active fragments of a processed oncogene protein  
 CC fragment as presented by a cancer cell or other antigen-presenting cell  
 CC and are capable of inducing a specific T-cell response to the actual  
 CC oncogene protein fragment as produced by the cell and processed and  
 CC presented in the HLA mol.  
 CC For example, a peptide fragment of p53 comprising at least  
 CC mutations in position 273, in which position any amino acid except Arg  
 CC may be located. The p53 sequence below was not disclosed in the  
 CC specification, but retrieved by the Indexer from Swiss-prot P04637.  
 SQ Sequence 393 AA;

Query Match 21.4%; Score 724.5; DB 1; Length 393;  
 Best Local Similarity 42.1%; Pred No. 2.8e-55;  
 Matches 160; Conservative 58; Mismatches 103; Indels 59; Gaps 10;  
 QY 14 TFEHLMSLEPSTYFDLPQSRGNEVYGGTSSMDVPHLEGMTTSVMAQFNLLSSTMD 73  
 DB 18 TTSDLKLLPENLVSLPLP--SQAMDDLMLSPD-----IEQWFTEDPGPD 61  
 QY 74 QMSRAASAPTYPEHAASVPHSPYAPQSSFTDMSAP-----VIPSNTDYPGPHF 127  
 DB 62 EAPRMEAPPVAPAPAAFTP-----AAP-----APAPSWPLSSVPSQTYOGSYGF 109  
 QY 128 EYTFQOSTAKSATWTYSPLLKLYCQIAKTCPIQIKVSTPPPPGTATAMPVYKKAHV 187  
 DB 110 RLGFHSGTAKSVTCYSPALNKMFCQLAKTCFQVLWVDSPTPPGTRVRAAIYKQSQM 169  
 QY 188 TDVVKPCPNHELGRDNQGSAPASHLRVEGNLNSQYVDDPVTGRQSVVVPYEPQVGT 247  
 DB 170 TEVVRCPHERCD-SDG-LAPPOHLIRVEGNLRVEYLDNRNTRFHSVVVPEPEVGS 227  
 QY 248 EETILYNFNCSSCVGNNRPILITILEMRDGOVLGRSFEGRICACGRDKADED 307  
 DB 228 DCTIHYNYCNSCMGNNRPILITILEDSGNNLLGRNSFEVRCACGRDRRTTEE 287  
 QY 308 HYREQOALNESSAKNG-----AASKRAFKQSPAPVAPALGAGYKRRHGDEDTYYLQV 359  
 DB 288 NLR-----KCEPHHELPPGSKRALPNNTSSSQ-----PKKPLDGEYFTLOI 332  
 QY 360 RGRNFELMKLESLELME 379  
 DB 333 RGRERFEMFRELNEALELKD 352

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